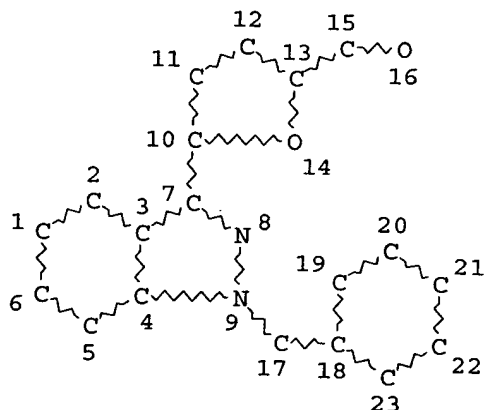


=> d que stat 123
L1 STR



NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L3 139 SEA FILE=REGISTRY SSS FUL L1
L21 1 SEA FILE=REGISTRY ABB=ON PLU=ON 170632-47-0
L22 138 SEA FILE=REGISTRY ABB=ON PLU=ON L3 NOT L21
L23 36 SEA FILE=HCAPLUS ABB=ON PLU=ON L22

=> d l23 ibib abs hitstr 1-36

L23 ANSWER 1 OF 36 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2005:1341973 HCAPLUS
TITLE: The design and synthesis of YC-1 analogs as probes for
soluble guanylate cyclase
AUTHOR(S): Hering, Kirk W.; Artz, Jennifer D.; Pearson, William
H.; Marletta, Michael A.
CORPORATE SOURCE: Department of Chemistry, University of Michigan, Ann
Arbor, MI, 48109-1055, USA
SOURCE: Bioorganic & Medicinal Chemistry Letters (2006),
16(3), 618-621
CODEN: BMCLE8; ISSN: 0960-894X
PUBLISHER: Elsevier B.V.
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Soluble guanylate cyclase (sGC) is highly activated in the presence of both
YC-1 [1-benzyl-3-(5-hydroxymethyl-2-furyl)indazole] and CO. The design,
synthesis, and activity (i.e., sGC activation) of photolabile analogs of
YC-1 are presented. Initial results with 6-azido-3-(5-hydroxymethyl-2-
furyl)-1-benzylindazole led to the synthesis of a tritium-labeled analog.
When photoactivated, this analog labeled the α -subunit of sGC.
IT 876365-94-5P
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic
preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant)

=> d his ful

(FILE 'HOME' ENTERED AT 14:15:25 ON 13 MAR 2006)

FILE 'REGISTRY' ENTERED AT 14:16:41 ON 13 MAR 2006

L1 STR
L2 10 SEA SSS SAM L1
L3 139 SEA SSS FUL L1

FILE 'HCAPLUS' ENTERED AT 14:18:20 ON 13 MAR 2006

L4 170 SEA ABB=ON PLU=ON L3

FILE 'REGISTRY' ENTERED AT 14:19:11 ON 13 MAR 2006

L5 STR L1
L6 121 SEA SUB=L3 SSS FUL L5

FILE 'HCAPLUS' ENTERED AT 14:20:07 ON 13 MAR 2006

L7 170 SEA ABB=ON PLU=ON L6

FILE 'REGISTRY' ENTERED AT 14:20:12 ON 13 MAR 2006

L8 STR L5
L9 0 SEA SUB=L3 SSS SAM L8
L10 1 SEA SUB=L3 SSS FUL L8
D SCA
L11 STR L8
L12 93 SEA SUB=L3 SSS FUL L11

FILE 'HCAPLUS' ENTERED AT 14:23:32 ON 13 MAR 2006

L13 170 SEA ABB=ON PLU=ON L12

FILE 'BEILSTEIN' ENTERED AT 14:24:50 ON 13 MAR 2006

L14 0 SEA SSS SAM L8
L15 0 SEA SSS FUL L8

FILE 'MARPAT' ENTERED AT 14:25:13 ON 13 MAR 2006

L16 0 SEA SSS SAM L8
L17 2 SEA SSS FUL L8

FILE 'HCAPLUS' ENTERED AT 14:25:30 ON 13 MAR 2006

L18 1 SEA ABB=ON PLU=ON L10

FILE 'MARPAT' ENTERED AT 14:25:34 ON 13 MAR 2006

L19 1 SEA ABB=ON PLU=ON L17 NOT L18

FILE HOME

FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 12 MAR 2006 HIGHEST RN 876514-29-3

DICTIONARY FILE UPDATES: 12 MAR 2006 HIGHEST RN 876514-29-3

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TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

FILE HCAPLUS

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FILE COVERS 1907 - 13 Mar 2006 VOL 144 ISS 12
FILE LAST UPDATED: 12 Mar 2006 (20060312/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

FILE BEILSTEIN

FILE LAST UPDATED ON JANUARY 17, 2006

FILE COVERS 1771 TO 2005.

FILE CONTAINS 9,428,406 SUBSTANCES

>>>PLEASE NOTE: Reaction Data and substance data are stored in separate documents and can not be searched together in one query. Reaction data for BEILSTEIN compounds may be displayed immediately with the display codes PRE (preparations) and REA (reactions). A substance answer set retrieved after the search for a chemical name, a compounds with available reaction information by combining with PRE/FA, REA/FA or more generally with RX/FA. The BEILSTEIN Registry Number (BRN) is the link between a BEILSTEIN compound and belonging reactions. For more detailed reaction searches BRNs can be searched as reaction partner BRNs Reactant BRN (RX.RBRN) or Product BRN (RX.PBRN).<<<

>>> FOR SEARCHING PREPARATIONS SEE HELP PRE <<<

* PLEASE NOTE THAT THERE ARE NO FORMATS FREE OF COST. *
* SET NOTICE FEATURE: THE COST ESTIMATES CALCULATED FOR SET NOTICE *
* ARE BASED ON THE HIGHEST PRICE CATEGORY. THEREFORE; THESE *
* ESTIMATES MAY NOT REFLECT THE ACTUAL COSTS. *
* FOR PRICE INFORMATION SEE HELP COST *

NEW

* PATENT NUMBERS (PN) AND BABS ACCESSION NUMBERS (BABSAN) CAN NOW BE
SEARCHED, SELECTED AND TRANSFERRED.
* NEW DISPLAY FORMATS ALLREF, ALLP AND BABSAN SHOW ALL REFERENCES,
ALL PATENT REFERENCES, OR ALL BABS ACCESSION NUMBERS FOR A
COMPOUND AT A GLANCE.

FILE MARPAT

FILE CONTENT: 1910-PRESENT VOL 144 ISS 11 (20060310/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1910-1987

MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
(COVERAGE TO THESE DATES IS NOT COMPLETE):

US	2006030554	09	FEB	2006
DE	102004053311	05	JAN	2006
EP	1609846	28	DEC	2005
JP	2006003337	05	JAN	2006
WO	2006012333	02	FEB	2006
GB	2415429	28	DEC	2005
FR	2873371	27	JAN	2006
RU	2266908	27	DEC	2005
CA	2495134	23	DEC	2005

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FILE CONTENT: 1910-PRESENT VOL 144 ISS 11 (20060310/ED)

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MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
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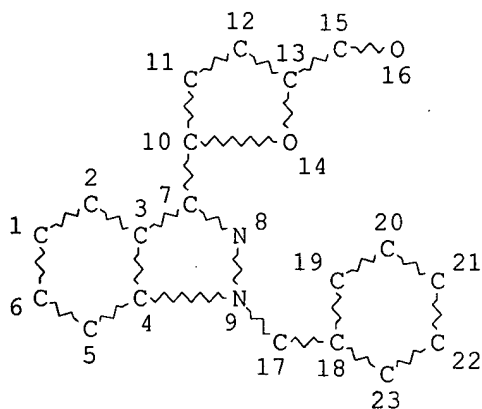
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EP	1609846	28	DEC	2005
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WO	2006012333	02	FEB	2006
GB	2415429	28	DEC	2005

FR 2873371 27 JAN 2006
 RU 2266908 27 DEC 2005
 CA 2495134 23 DEC 2005

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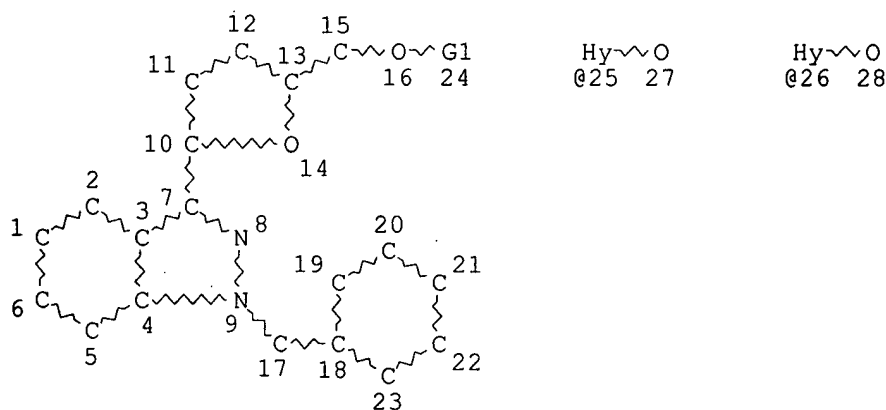
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 L1 STR



NODE ATTRIBUTES:
 DEFAULT MLEVEL IS ATOM
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
 RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE
 L3 139 SEA FILE=REGISTRY SSS FUL L1
 L8 STR



VAR G1=25/26
 NODE ATTRIBUTES:
 DEFAULT MLEVEL IS ATOM

GGCAT IS MCY SAT AT 25
GGCAT IS MCY SAT AT 26
DEFAULT ECLEVEL IS LIMITED
ECOUNT IS E5 C E1 O AT 25
ECOUNT IS E4 C E1 O AT 26

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 28

STEREO ATTRIBUTES: NONE
L10 1 SEA FILE=REGISTRY SUB=L3 SSS FUL L8
L18 1 SEA FILE=HCAPLUS ABB=ON PLU=ON L10

=> d l18 ibib abs hitstr
YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L18 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2004:902206 HCAPLUS
DOCUMENT NUMBER: 141:388641
TITLE: HIF-1 expression inhibition-based method for
inhibiting tumor angiogenesis and tumor growth
INVENTOR(S): Park, Jong-Wan; Chun, Yang-Sook; Kim, Jinho
PATENT ASSIGNEE(S): Bizbiotech Co., Ltd., USA
SOURCE: PCT Int. Appl., 69 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004091648	A1	20041028	WO 2004-US10327	20040401
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004198798	A1	20041007	US 2003-407136	20030407
US 2005096370	A1	20050505	US 2003-642363	20030814
PRIORITY APPLN. INFO.:			US 2003-407136	A 20030407
			US 2003-642363	A 20030814

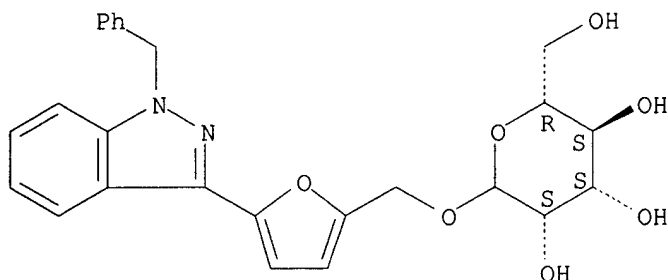
OTHER SOURCE(S): MARPAT 141:388641
AB The invention provides methods and pharmaceutical compns. for inhibiting expression of HIF-1 and HIF-1-regulated genes, angiogenesis, tumor growth, or tumor progression/metastasis, comprising contacting the tumor cells or tissue with a composition comprising 3-(5'-hydroxymethyl-2'-furyl)-1-benzylindazole, or a derivative thereof.
IT 781663-33-0

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
(HIF-1 expression inhibition-based method for inhibiting tumor
angiogenesis and tumor growth)

RN 781663-33-0 HCAPLUS

CN D-Mannopyranoside, [5-[1-(phenylmethyl)-1H-indazol-3-yl]-2-furanyl]methyl
(9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> fil marpat

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(COVERAGE TO THESE DATES IS NOT COMPLETE):

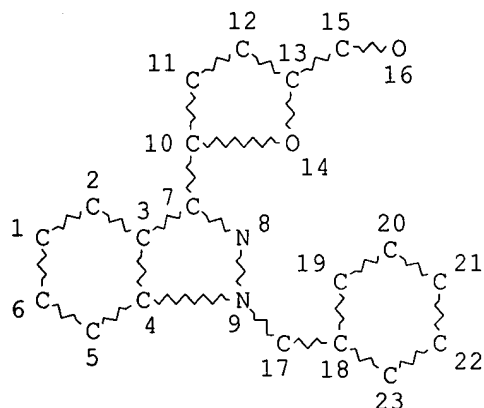
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DE	102004053311	05	JAN	2006
EP	1609846	28	DEC	2005
JP	2006003337	05	JAN	2006
WO	2006012333	02	FEB	2006
GB	2415429	28	DEC	2005
FR	2873371	27	JAN	2006
RU	2266908	27	DEC	2005
CA	2495134	23	DEC	2005

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=> d que stat 119

L1 STR



NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

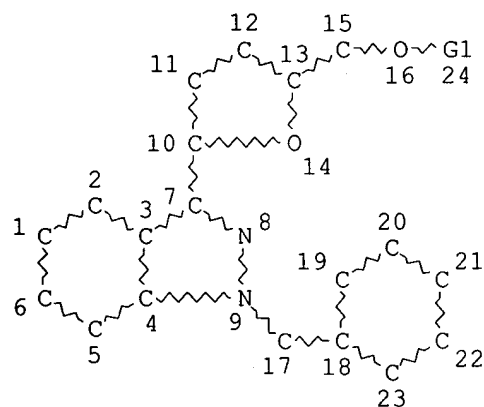
RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L3 139 SEA FILE=REGISTRY SSS FUL L1

L8 STR



VAR G1=25/26

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

GGCAT IS MCY SAT AT 25

GGCAT IS MCY SAT AT 26

DEFAULT ECLEVEL IS LIMITED

ECOUNT IS E5 C E1 O AT 25

ECOUNT IS E4 C E1 O AT 26

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 28

STEREO ATTRIBUTES: NONE

L10 1 SEA FILE=REGISTRY SUB=L3 SSS FUL L8
 L17 2 SEA FILE=MARPAT SSS FUL L8
 L18 1 SEA FILE=HCAPLUS ABB=ON PLU=ON L10
 L19 1 SEA FILE=MARPAT ABB=ON PLU=ON L17 NOT L18

=> d 119 ibib abs qhit

L19 ANSWER 1 OF 1 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 142:441845 MARPAT
 TITLE: Method for inhibiting tumor angiogenesis and tumor growth
 INVENTOR(S): Park, Jong-Wan; Chun, Yang-Sook; Kim, Jinho
 PATENT ASSIGNEE(S): Bizbiotech Co., Ltd., S. Korea
 SOURCE: U.S. Pat. Appl. Publ., 36 pp., Cont.-in-part of U.S. Ser. No. 407,136.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

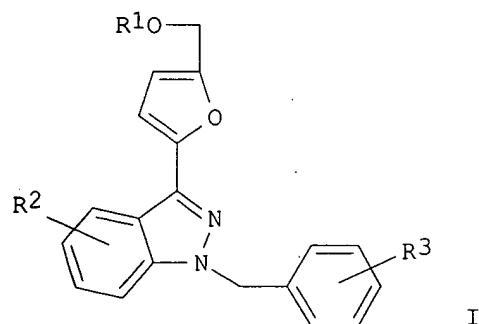
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005096370	A1	20050505	US 2003-642363	20030814
US 2004198798	A1	20041007	US 2003-407136	20030407
WO 2004091648	A1	20041028	WO 2004-US10327	20040401

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

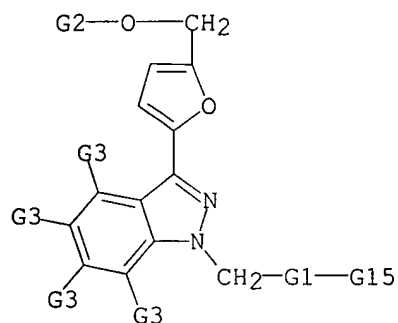
PRIORITY APPLN. INFO.: US 2003-407136 20030407
 US 2003-642363 20030814

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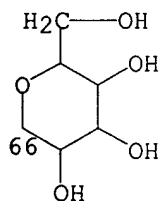


AB The present invention provides methods and pharmaceutical compns. for

inhibiting expressions of HIF-1 and HIF-1-regulated genes, angiogenesis, tumor growth, or tumor progression/metastasis comprising contacting the tumor cells or tissue with a composition comprising 3-(5'-hydroxymethyl-2'-furyl)-1-benzylindazole, or mixture of compds. of the formula I (where R1 = polyol, R2, R3 = H, alkyl, alkoxy, halogen, etc.).

MSTR 1

G1 = phenylene
G2 = 66



Patent location:

Note:

Note:

Note:

Stereochemistry:

claim 7

substitution is restricted

additional ring and oxo formation also disclosed

and pharmaceutically acceptable solvates and salts

and isomers and mixtures of isomers